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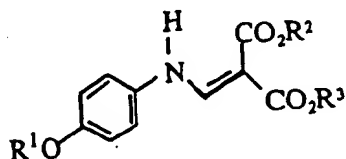
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(54) Cosmetic light-screening composition

(57) The invention relates to a cosmetic light-screening composition for protecting human skin or human hair against ultraviolet radiation containing at least one compound of formula I



I

wherein

R¹ represents a C₁₋₈ straight or branched alkyl chain, and
R² and R³ each independently represent a C₁₋₈ straight or branched alkyl chain.

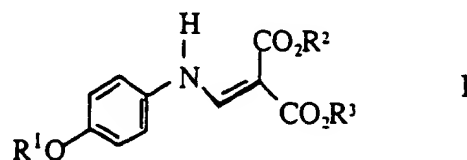
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Description

[0001] The invention relates to a cosmetic light-screening composition, the use of the cosmetic light-screening composition for the protection of the human skin and human hair against the ultraviolet radiation of wavelengths between about 280 and 400 nm, compounds contained in the composition and the use of these compounds as UV filters.

[0002] It is known that sunlight accelerates the ageing of skin and even gives rise to skin cancer, these undesired effects being caused by UV radiation.

[0003] It has now been found that the compounds of the formula I



wherein

R¹ represents a C₁₋₈ straight or branched alkyl chain, and

R² and R³ each independently represent a C₁₋₈ straight or branched alkyl chain are excellent UV filters concerning skin compatibility and stability (light, heat, moisture); they make a strong contribution to the UV protection of the skin and therewith cause a delay in skin ageing. In particular these UV filters also have an outstanding photostability.

[0004] It has also been found that the compounds of formula I have their absorption maxima between those of UV A-filters like e.g. 4-tert-butyl-4'-methoxydibenzoylmethane and those of UV B filters and especially and surprisingly increase the protective effect of UV B filters, i.e. of substances which mainly absorb the erythema-producing UV-B radiation in the region of about 290 to about 320 nm, although the absorption maximum of the compound of formula I does not lie in this region, but in the region of about 320 to 340 nm, i.e. between the maxima of UVA and UVB radiation.

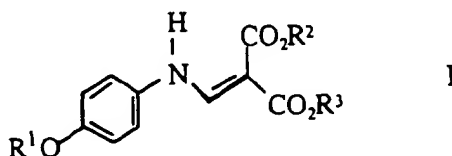
[0005] The only types of UV-filters available for cosmetic use which also absorb in these regions are the benzophenones and menthyl anthranilate. Many formulators don't use them for various reasons, e.g. very low extinction compared to the compounds of formula I.

[0006] Alkoxyanilinomethylene-propanedioic acid esters of the formula I are known from US Patent No. 3,079,366 and from W.O. Kermack, N.E. Storey, J. Chem. Soc 607 (1950); T. Takahashi, S. Senda, J. Pharm. Soc. Jpn., 71, 1112 (1952). Specifically disclosed are methoxy-ethoxy- and butoxy-anilinomethylene-propanedioic ethyl esters. But nothing is disclosed about the use of these compounds for cosmetic purposes, especially to use these compounds in skin care or hair care.

[0007] An UV-filter should show a large solubility in cosmetic solvent in order to be formulated in a reasonable concentration. For instance, a 6% UV-filter content in a classical o/w emulsion having a 70% water content requires a solubility up to 20% in the oil phase. Surprisingly, it has been found that the compounds of formula I show solubility up to 20% in many cosmetic solvents like caprylic capric triglyceride, propylene glycol dicaprylate/dicaprate, C₁₂₋₁₅ alkyl benzoate, propylene glycol monoisostearate, diisopropyladipate.

[0008] Objects of the present invention are accordingly novel compounds of formula I, light screening preparations for cosmetic purposes containing at least one of the compounds of formula I above, preferably in combination with at least one UV B filter agent and at least one UV A filter agent, and the use of compounds of formula I as light screening agents, especially for cosmetic purposes.

[0009] According to the present invention there is provided a cosmetic light-screening composition for protecting human skin or human hair against ultraviolet radiation containing in a cosmetically acceptable carrier at least one compound of formula I



wherein

R¹ represents a C₁₋₈ straight or branched alkyl chain,
R² and R³ each independently represent a C₁₋₈ straight or branched alkyl chain.

[0010] The term "C₁₋₈ straight or branched alkyl chain" refers to groups like methyl, ethyl, n-propyl, isopropyl, n-butyl, tert.butyl, pentyl, heptyl, 2-ethylhexyl and the like.

[0011] Preferred are compounds wherein R² and R³ each independently represent methyl, ethyl, pentyl or 2-ethylhexyl.

[0012] The group R¹ is preferably methyl, ethyl or n-butyl.

[0013] Thus, the following compounds are preferred:

compounds wherein R¹, R² and R³ are methyl (Ex.1) or ethyl (Ex.4); or
R¹ is ethyl and R² and R³ are methyl (Ex.2) or 2-ethylhexyl (Ex.6); or
R¹ is n-butyl and R² and R³ are methyl. (Ex.3) or ethyl (Ex.5).

[0014] Most preferred is a compound according to Example 4 wherein R¹, R² and R³ are ethyl.

[0015] The cosmetic light screening composition comprises preferably 0.1 to 10% by weight, more preferably 0.5 to 5% by weight, of the compound(s) of formula I, in particular 1 to 3% by weight of the compound(s) of formula I.

[0016] The cosmetic light screening composition may in addition comprise at least one UV-B filter agent with an absorption maximum at about 300 to 320 nm and at least one UV-A filter agent with an absorption maximum at ≥ 340 nm, especially at between 340 and 360 nm, particularly one UV-A filter agent having an absorption maximum at 356 nm. Specifically this UV-A filter agent is 4-tert-butyl-4'-methoxydibenzoylmethane. Generally the UV-A filter agent can be chosen out of the group consisting of 2-methyldibenzoylmethane, 4-methyl-dibenzoyl-methane, 4-tert-butylidibenzoylmethane, 2,4-dimethyldibenzoylmethane, 2,5-dimethyldibenzoylmethane, 4,4'-diisopropyldibenzoylmethane, 4-tert-butyl-4'-methoxydibenzoylmethane, 2-methyl-5-isopropyl-4'-methoxydibenzoylmethane, 2-methyl-5-tert-butyl-4'-methoxydibenzoylmethane, 2,4-dimethyl-4'-methoxydibenzoylmethane and 2,6-dimethyl-4-tert-butyl-4'-methoxydibenzoylmethane and terephthalylidene dicamphor sulfonic acid.

[0017] The UV-B filter agent is at least one of the group consisting of cinnamates, salicylates, benzophenones, diphenylacrylates triazines, camphor derivatives, polymeric UV absorbers specially those described in US applications US 5 403 944 and microfine pigments. Specifically the UV-B filter agent is at least one of the group consisting of pigment metallic oxides of cerium, iron, titanium, zinc or zirconium, especially of titanium or zinc, and polymers with hydrocarbon structure or siloxane structure carrying at least one ultraviolet-light-absorbing group.

[0018] The cosmetic light-screening compositions or the compounds of formula I are useful for protecting human skin or human hair against ultraviolet radiation.

[0019] The following compounds of formula I are new:

2-(4-Heptoxy-anilinomethylene)-propanedioic acid diethyl ester,
2-(4-Methoxy-anilinomethylene)-propanedioic acid dimethyl ester,
2-(4-Ethoxy-anilinomethylene)-propanedioic acid dimethyl ester,
2-(4-Propoxy-anilinomethylene)-propanedioic acid dimethyl ester,
2-(4-Butoxy-anilinomethylene)-propanedioic acid dimethyl ester,
2-(4-Pentoxy-anilinomethylene)-propanedioic acid dimethyl ester,
2-(4-Hexoxy-anilinomethylene)-propanedioic acid dimethyl ester,
2-(4-Ethoxy-anilinomethylene)-propanedioic acid dipentyl ester,
2-(4-Ethoxy-anilinomethylene)-propanedioic acid ethyl ester pentyl ester, 2-(4-Ethoxy-anilinomethylene)-propanedioic acid di-2-ethyl-hexyl ester and 2-(4-Ethoxy-anilinomethylene)-propanedioic acid ethyl ester 2-ethyl-hexyl ester.

[0020] Each of those can be used as an UV filter, especially for cosmetic purposes as said already before. In this respect the following ones are of specific interest:

2-(4-Ethoxy-anilinomethylene)-propanedioic acid dipentyl ester,

5 2-(4-Ethoxy-anilinomethylene)-propanedioic acid ethyl ester pentyl ester, 2-(4-Ethoxy-anilinomethylene)-propanedioic acid di-2-ethyl-hexyl ester and 2-(4-Ethoxy-anilinomethylene)-propanedioic acid ethyl ester 2-ethyl-hexyl ester.

[0021] The compounds represented by formula I (known and new compounds) are conveniently prepared by conventional methods as disclosed in US 3,079,366 and which may be described briefly as follows:

[0022] Reaction of an aniline derivatives (C₁-C₈ alkoxyaniline) with the appropriately substituted alkoxyalkylene-malono derivative with or without a solvent.

[0023] As cosmetically acceptable carrier usual for light screening agents in the scope of the present invention there can be used any conventional preparation which corresponds to the cosmetic requirements, e.g. creams, lotions, emulsions, 15 salves, gels, solutions, sprays, sticks and milks; see e.g. Kosmetik, Entwicklung, Herstellung und Anwendung Kosmetischer Mittel, ed. Wilfried Umbach, Georg Thieme Verlag Stuttgart - New York 1988; Sunscreens, Development, Evaluation and Regulatory Aspects, ed. N.Y. Lowe, N.A. Shaat, Marcel Decker, Inc. New York Basel, 1990.

[0024] Having regard to their lipophilicity, the compounds of formula I can be incorporated well in oil-containing and fat-containing cosmetic preparations.

20 [0025] With respect to the lipophilicity, the novel compounds fulfil the criteria which are required in the present instance, namely a solubility in cosmetic solvents, such as e.g. Miglyol 812N (caprylic capric triglyceride), Miglyol 840 (propylene glycol dicaprylate dicaprate), Finsolv TN (C₁₂₋₁₅ alkyl benzoate), Prisorine 2034 (propylene glycol monoiso-stearate) or Crodamol DA (diisopropyladipate).

[0026] The compound of Example 4 is especially preferred particularly in the just mentioned respect.

25 [0027] Suitable the cosmetic screening formulations takes the form of an oil, a lotion, a gel, a solid stick, an emulsion, e.g. cream, milk or of a vesicular dispersion of ionic or nonionic amphiphilic lipids, an aerosol, a spray, a foam, a powder, a shampoo, a hair conditioner or lacquer or a make-up or the like.

[0028] In case a cosmetic formulation for the protection of human hair is prepared by at least one compound of formula I the suitable formulations are shampoos, conditioners, lotions, gels, emulsions, dispersions, lacquers or the like.

30 The preparation of all these formulations is well known to the skilled artisan.

[0029] The usual solvents known to the skilled artisan can be used for the preparation of these forms, e.g. oils, waxes, alcohols, polyols. The preferred agents are fatty acids, esters, fatty alcohols, but also ethanol, isopropanol, propylene glycol, glycerine or the like are useful.

35 [0030] The cosmetic formulations may contain further adjuvants, e.g. further solvents, thickeners, emollients, emulsifiers, humectants, tensides, preservatives, antifoams, fragrances, oils, waxes, lower polyols and monohydric alcohols, propellants, silicones, colourings and pigments or the like.

[0031] An important advantage of the novel light-screening composition or the compounds of formula I stems from the fact that the artisan skilled in the art is completely free in the choice regarding the material used for the filtration of the UV-B and UV-A radiation as already said above. But for UV-A filtration the most preferred UV-A filter agent is 4-tert-butyl-4'-methoxydibenzoylmethane, this is especially used in combination with 2-(4-ethoxy-anilinomethylene)-propanedioic acid diethyl ester.

[0032] Further advantages, characteristics and details are disclosed by the following examples.

Example 1

45 2-(4-Methoxy-anilinomethylene)-propanedioic acid dimethyl ester

[0033] To a solution of 12.3 g (0.1 Mol) of p-Anisidine in 50 ml of ethanol, a solution of 17.4 g (0.1 Mol) of Dimethyl methoxymethylenemalonate in 70 ml of ethanol was added dropwise with stirring at room temperature. After an additional 50 hour of stirring at that temperature, the reaction mixture was cooled with an ice bath and the product crystallised spontaneously. The crude solid material was isolated by filtration and washed with ethanol and gave 12.9 g of the title product which is a white solid melting at 89-91° C, UV 328 nm (E=899).

Example 2

55 2-(4-Ethoxy-anilinomethylene)-propanedioic acid dimethyl ester

[0034] Same preparation as in example 1 where one equivalent of p-Phenetidine was used instead of p-Anisidine.

19.8 g of the title compound was obtained which is a white solid melting at 66-67 °C, UV 328 nm (E=879).

Example 3

2-(4-Butoxy-anilinomethylene)-propanedioic acid dimethyl ester

[0035]

- a) Preparation of 2-(4-Hydroxy-anilinomethylene)-propanedioic acid dimethyl ester.
 To a solution of 16.4 g (0.15 Mol) of 4-Aminophenol in 100 ml of ethanol, a solution of 26.1 g (0.15 Mol) of Dimethyl methoxymethylenemalonate in 70 ml of ethanol was added dropwise at room temperature. After an additional 10 minutes of stirring at that temperature the crude solid material was isolated by filtration and washed with ethanol and gave 34.3g of a white solid which was then used without further purification.
- b) To a mixture of 7 g (28 mMol) of 2-(4-Hydroxy-anilinomethylene)propanedioic acid dimethyl ester, 3.9 g (28 mMol) of Potassium carbonate in 50 ml of dimethylformamide, 7.3 g (28 mMol) of Butyl iodide were added dropwise with stirring at room temperature. The reaction mixture was heated at 80 °C for 24 hours. Then the reaction mixture was poured into 50 ml of water and extracted three times with ethyl acetate. The organic layer was washed with a solution of sodium hydroxide (15%) and thereafter with water, dried and evaporated and gave 4.5 g of the title compound which is a white solid melting at 46-47 °C, UV 328 nm (E=685).

Example 4

2-(4-Ethoxy-anilinomethylene)-propanedioic acid diethyl ester

- The preparation was the same as in example 1. But one equivalent of p-Phenetidine was used instead of p-Anisidine and one equivalent of diethyl ethoxymethylenemalonate was used instead of dimethyl methoxymethylenemalonate. Ethanol was replaced by hexane. 21 g of the title compound was obtained which is a white solid melting at 55-56 °C, UV 329 nm (E=802).

Example 5

2-(4-Butoxy-anilinomethylene)-propanedioic acid diethyl ester

- The preparation was the same as in example 3. But one equivalent of diethyl ethoxymethylenemalonate was used instead of dimethyl methoxymethylenemalonate. 3.1 g of the title compound was obtained which is a white solid melting at 54-55 °C, UV 329 nm (E=714).

Example 6

2-(4-Ethoxy-anilinomethylene)-propanedioic acid di-2-ethyl-hexyl ester and 2-(4-Ethoxy-anilinomethylene)-propanedioic acid ethyl ester 2-ethyl-hexyl ester

- [0036] A mixture of 18.4 g (60 mMol) of 2-(4-Ethoxy-anilinomethylene)propanedioic acid diethyl ester, 430 mg of titanium(IV) isopropoxide in 70 ml of 2-Ethyl-hexanol was stirred at 150 °C for 4 hours. The reaction mixture was concentrated under reduced pressure (80 °C, 6 mbar) to get a yellow oil. This oil was dissolved into 100 ml of ethyl acetate and 5 ml of water, stirred for 10 min and dried over anhydrous magnesium sulfate. The flaky suspension was filtered and concentrated under reduced pressure and gave 26.8 g of an yellow oil. The two title products were isolated by flash chromatography using hexane: ethyl acetate 10:1 as eluant.

- [0037] 23.6 g of 2-(4-Ethoxy-anilinomethylene)-propanedioic acid di-2-ethylhexyl ester was obtained which is a yellow oil, UV 329 nm (E=567).

- [0038] 1.8 g of Ethoxy-anilinomethylene)-propanedioic acid ethyl ester and 2-ethyl-hexyl ester which is a yellow oil UV 329 nm (E=632).

Example 7

2-(4-Ethoxy-anilinomethylene)-propanedioic acid dipentyl ester and 2-(4-Ethoxy-anilinomethylene)-propanedioic acid ethyl ester pentyl ester

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[0039] The preparations were the same as in example 6 where n-pentanol was used instead of 2-Ethyl-hexanol. The two title products were isolated by flash chromatography using hexane: ethyl acetate 8:1 as eluant.

[0040] 16.6 g of 2-(4-Ethoxy-anilinomethylene)-propanedioic acid dipentyl ester was obtained which is a yellow oil, UV 329 nm (E=651).

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[0041] 2.5 g of Ethoxy-anilinomethylene)-propanedioic acid ethyl ester pentyl ester was obtained which is a yellow oil UV 329 nm (E=725).

Example 8

15 Preparation of formula type: water in oil emulsion

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	Ingredients	% w/w
5	A)	
	Product from example 6 (di-2-ethyl-hexyl ester)	6
10	Butyl Methoxydibenzoylmethane (Parsol 1789)	1.5
	Octyl methoxycinnamate (Parsol MCX)	3
15	Polyglyceryl-3 Diisostearate	5
	Glyceryl oleate	3
20	Cetearyl alcohol	2
	Mineral oil	10
25	Coco caprylate/caprate	10
	Titanium Dioxide coated with Dimethicone	2
30	Octyldodecanol	2
	Butylhydroxytoluene	0.1
35	Phenoxyethanol and Methylparaben and	0.6
	Ethylparaben and Propylparaben and Butylparaben	
40	B)	
	Glycerol (86%)	5
45	Phenylbenzimidazole Sulfonic Acid (Parsol HS)	2
	Disodium EDTA	0.1
50	Water	47.7

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[0042] Part A and part B were mixed separately at 85°C and then combined under stirring. Finally, the pH was corrected to 7 with potassium hydroxide 10% or citric acid 10% if necessary.

Example 9

Preparation of formula type: water in oil emulsion (soft cream)

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Ingredients	% w/w
A)	
Product from example 4	5
POP-POE Glycerol sorbitan fatty acids Esters	5
Heptamethylnonane	5
Paraffin	9
PPG-(15)-stearyl alcohol and cyclomethicone	2
silica	0.4
Butylhydroxytoluene	0.1
Phenoxyethanol and Methylparaben and	0.6
Ethylparaben and Propylparaben and Butylparaben	
B)	
Sodium chloride	0.5
POE-30 Sorbitol	1.5
Glycerol (86%)	2.5
Disodium EDTA	0.1
Water	68.3

Part A and part B were mixed separately at 85°C and then combined under stirring. Finally, the pH was corrected to 7
 55 with potassium hydroxide 10% or citric acid 10% if necessary.

Example 10

Formula type: silicon in water emulsion (lotion)

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Ingredients	% w/w
A)	
Product from example 5	2
Cyclomethicone pentamere and	9
Aluminium magnesium hydroxy stearate	
Cyclomethicone and Dimethicone Copolyol	10
Cyclomethicone	5
PPG-3 Myristyl ether	2
Titanium dioxide coated with Dimethicone	2
C12-15 Alkylbenzoate	10
Butylhydroxytoluene	0.1
Phenoxyethanol and Methylparaben and	0.6
Ethylparaben and Propylparaben and Butylparaben	
B)	
Sodium chloride	0.5
Tocopherylacetate	2
Disodium EDTA	0.1
Water	56.7

Part A and part B were mixed separately at 85°C and then combined under stirring. Finally, the pH was corrected to 7 with potassium hydroxide 10% or citric acid 10% if necessary.

Example 11

Preparation of formula type: oil in water emulsion

Ingredients	% w/w
A)	
Product from example 4	4
Octyl methoxycinnamate (Parsol MCX)	3
Butyl methoxydibenzoylmethane (Parsol 1789)	1.5
4-Methylbenzilidene Camphor (Parsol 5000)	3
Glyceryl Monomyristate	4
Cetyl alcohol	1
Coco caprilate caprate	15
Isopropyl myristate	5
PVP-Eicosen copolymer	2
Butylhydroxytoluene	0.1
Disodium EDTA	0.1
Phenoxyethanol and Methylparaben and	0.6
Ethylparaben and Propylparaben and Butylparaben	
B)	
POE-POP Block copolymer	2
Water	38.7

5	Carbomer 981	10
	Propylene glycol	10

- 10 Part A and part B were mixed separately at 85°C and then combined under stirring. Finally, the pH was corrected to 7 with potassium hydroxide 10% or citric acid 10% if necessary.

Example 12

- 15 Formula type: Oil in water emulsion

20	Ingredients	% w/w
	A)	
25	Product from example 4	3.5
	Octyl methoxycinnamate (Parsol MCX)	3
30	Butyl methoxydibenzoylmethane (Parsol 1789)	1.5
	4-Methylbenzilidene camphor (Parsol 5000)	3
35	Glyceryl monomyristate	4
	Cetyl alcohol	1
40	Coco caprilate caprate	5
	Caprilic capric triglyceride	5
45	PVP-Eicosen copolymer	1
	Butylhydroxytoluene	0.1
50	Disodium EDTA	0.1
	Phenoxyethanol and Methylparaben and	0.6
55	Ethylparaben and Propylparaben and Butylparaben	

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B)

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Sorbitan ester and Sucrose ester

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Water

48.2

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Carbomer 981

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Propylene glycol

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[0043] Part A and part B were mixed separately at 85°C and then combined under stirring. Finally, the pH was corrected to 7 with potassium hydroxide 10% or citric acid 10% if necessary.

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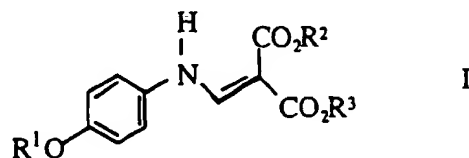
[0044] The emulsions of Examples 8, 11 and 12 showed a broad filter activity over the complete UV region of about 280 to 400 nm with a broad plateau maximum in the range of about 300 to 360 nm.

Claims

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1. A cosmetic light-screening composition for protecting human skin or human hair against ultraviolet radiation containing in a cosmetically acceptable carrier at least one compound of formula I

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wherein

R¹ represents a C₁₋₈ straight or branched alkyl chain, and
R² and R³ each independently represent a C₁₋₈ straight or branched alkyl chain.

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2. A cosmetic light-screening composition according to claim 1 wherein

R¹ represents methyl, ethyl or n-butyl, and
R² and R³ each independently represents methyl, ethyl, pentyl, 2-ethylhexyl.

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3. A cosmetic light-screening composition according to claim 2 wherein R¹, R² and R³ are ethyl.

4. A cosmetic light-screening composition according to any one of claims 1 to 3 comprising 0.1 to 10% by weight, preferably 0.5 to 5% by weight of the compound(s) of formula I.

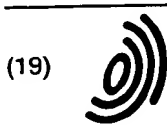
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5. A cosmetic light-screening composition according to claim 4 comprising 1 to 3% by weight of the compound(s) of formula I.

6. A cosmetic light-screening composition according to any one of claims 1 to 5 comprising in addition at least one

UV-B filter agent with an absorption maximum at about 300 to 320 nm and at least one UV-A filter agent with an absorption maximum at ≥ 340 nm, especially between 340 and 360 nm.

- 5 7. A cosmetic light-screening composition according to claim 6 wherein the UV-A filter agent is one of the group consisting of 2-methyldibenzoylmethane, 4-methyl-dibenzoyl-methane, 4-tert-butyldibenzoyl-methane, 2,4-dimethyldibenzoylmethane, 2,5-dimethyldibenzoylmethane, 4,4'-diisopropyldibenzoylmethane, 4-tert-butyl-4'-methoxydibenzoylmethane, 2-methyl-5-isopropyl-4'-methoxydibenzoylmethane, 2-methyl-5-tert-butyl-4'-methoxydibenzoylmethane, 2,4-dimethyl-4'-methoxydibenzoylmethane, 2,6-dimethyl-4-tert-butyl-4'-methoxydibenzoylmethane and terephthalylidene dicamphor sulfonic acid.
- 10 8. A cosmetic light-screening composition according to claim 7 wherein the UV-A filter agent is 4-tert-butyl-4'-methoxydibenzoylmethane.
- 15 9. A cosmetic light-screening composition according to any one of claims 6 to 8 wherein the UV-B filter agent is at least one of the group consisting of cinnamates, salicylates, benzophenones, diphenylacrylates triazines, camphor derivatives, polymeric UV-absorbers or pigment metallic oxides of cerium, iron, titanium, zinc or zirconium, especially of titanium or zinc, and polymers with hydrocarbon structure or siloxane structure carrying at least one ultraviolet-light-absorbing group.
- 20 10. Use of a cosmetic light-screening composition of any one of the claims 1 to 9 for protecting human skin or human hair against ultraviolet radiation.
11. Compounds of formula I selected from the group consisting of
 - 25 2-(4-Heptoxy-anilinomethylene)-propanedioic acid diethyl ester,
 - 2-(4-Methoxy-anilinomethylene)-propanedioic acid dimethyl ester,
 - 2-(4-Ethoxy-anilinomethylene)-propanedioic acid dimethyl ester,
 - 2-(4-Propoxy-anilinomethylene)-propanedioic acid dimethyl ester,
 - 30 2-(4-Butoxy-anilinomethylene)-propanedioic acid dimethyl ester,
 - 2-(4-Pentoxy-anilinomethylene)-propanedioic acid dimethyl ester,
 - 2-(4-Hexoxy-anilinomethylene)-propanedioic acid dimethyl ester,
 - 2-(4-Ethoxy-anilinomethylene)-propanedioic acid dipentyl ester,
 - 2-(4-Ethoxy-anilinomethylene)-propanedioic acid ethyl ester pentyl ester, 2-(4-Ethoxy-anilinomethylene)-propanedioic acid di-2-ethyl-hexyl ester and 2-(4-Ethoxy-anilinomethylene)-propanedioic acid ethyl ester 2-ethyl-hexyl ester.
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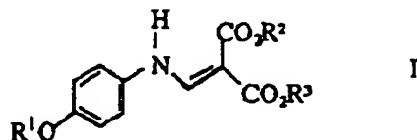
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(54) Cosmetic light-screening composition

(57) The invention relates to a cosmetic light-screening composition for protecting human skin or human hair against ultraviolet radiation containing at least one compound of formula I



wherein

R¹ represents a C₁₋₈ straight or branched alkyl chain, and

R² and R³ each independently represent a C₁₋₈ straight or branched alkyl chain.

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European Patent
Office

EUROPEAN SEARCH REPORT

Application Number
EP 98 11 4262

DOCUMENTS CONSIDERED TO BE RELEVANT			
Category	Citation of document with indication, where appropriate, of relevant passages	Relevant to claim	CLASSIFICATION OF THE APPLICATION (Int.Cl.6)
D,X	US 3 079 366 A (R.J.BOYLE ET AL) 26 February 1963 * column 2, line 5 - line 30; claim 1; table IV *	1	A61K7/42
A	US 5 639 446 A (G. RASPANTI ET AL) 17 June 1997 * claims 1,2 *	1	
A	DE 41 22 475 A (BASF) 7 January 1993 * claims 1,9,12 *	1	
A	WO 96 15102 A (BASF) 23 May 1996 * claims 1,12 *	1	
A	EP 0 174 832 A (MAY & BAKER) 19 March 1986 * page 55, line 5 *	11	
A	FR 2 002 888 A (WARNER-LAMBERT) 31 October 1969 * table I *	11	
The present search report has been drawn up for all claims			TECHNICAL FIELDS SEARCHED (Int.Cl.6)
			A61K
Place of search	Date of completion of the search	Examiner	
THE HAGUE	10 March 1999	VOYIAZOGLU D.	
CATEGORY OF CITED DOCUMENTS		T : theory or principle underlying the invention E : earlier patent document, but published on, or after the filing date D : document cited in the application L : document cited for other reasons & : member of the same patent family, corresponding document	
X : particularly relevant if taken alone Y : particularly relevant if combined with another document of the same category A : technological background O : non-written disclosure P : intermediate document			

EPO FORM 1503 03.82 (P04C01)

**ANNEX TO THE EUROPEAN SEARCH REPORT
ON EUROPEAN PATENT APPLICATION NO.**

EP 98 11 4262

This annex lists the patent family members relating to the patent documents cited in the above-mentioned European search report.
The members are as contained in the European Patent Office EDP file on
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10-03-1999

Patent document cited in search report		Publication date	Patent family member(s)	Publication date
US 3079366	A	26-02-1963	FR 1330982 A NL 279734 A	16-12-1963
US 5639446	A	17-06-1997	NONE	
DE 4122475	A	07-01-1993	AT 128119 T CA 2109594 A DE 59203777 D WO 9301164 A EP 0593541 A JP 6509077 T SK 150193 A US 5443820 A	15-10-1995 21-01-1993 26-10-1995 21-01-1993 27-04-1994 13-10-1994 06-07-1994 22-08-1995
WO 9615102	A	23-05-1996	DE 4440055 A DE 19519895 A AU 3980395 A BG 101460 A BR 9509644 A CZ 9701349 A EP 0790980 A EP 0900782 A FI 971991 A JP 10511081 T NO 972156 A PL 320203 A SK 56097 A US 5821380 A	15-05-1996 05-12-1996 06-06-1996 31-08-1998 16-09-1997 15-04-1998 27-08-1997 10-03-1999 09-05-1997 27-10-1998 09-07-1997 15-09-1997 08-07-1998 13-10-1998
EP 174832	A	19-03-1986	JP 61087653 A US 4659727 A	06-05-1986 21-04-1987
FR 2002888	A	31-10-1969	DE 1908548 A GB 1237059 A	05-11-1970 30-06-1971

EPO FORM P0458

For more details about this annex : see Official Journal of the European Patent Office, No. 12/82

